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SALES hereby certify that annexed is a true copy of the Provisional specification
in connection with Application No. 2002952129 for a patent by THE
UNIVERSITY OF QUEENSLAND as filed on 17 October 2002.



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Sixth day of November 2003

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PROVISIONAL SPECIFICATION

Applicant:

THE UNIVERSITY OF QUEENSLAND

Invention Title:

TREATMENT OF HYPERSENSITIVITY CONDITIONS

The invention is described in the following statement:

TREATMENT OF HYPERSENSITIVITY CONDITIONS

FIELD OF THE INVENTION

5 This invention relates to novel cyclic compounds
which have the ability to modulate the activity of G
protein-coupled receptors. The compounds preferably act
as antagonists. In preferred embodiments, the invention
provides cyclic peptidic and peptidomimetic antagonists of
C5a receptor, which are active against C5a receptors on
10 polymorphonuclear leukocytes and macrophages. The
compounds of the invention are both potent and selective,
and are useful in the treatment of hypersensitivity
conditions such as asthma and other allergic conditions.

15 BACKGROUND OF THE INVENTION

All references, including any patents or patent
applications, cited in this specification are hereby
incorporated by reference. No admission is made that any
reference constitutes prior art. The discussion of the
20 references states what their authors assert, and the
applicants reserve the right to challenge the accuracy and
pertinency of the cited documents. It will be clearly
understood that, although a number of prior art
publications are referred to herein, this reference does
25 not constitute an admission that any of these documents
forms part of the common general knowledge in the art, in
Australia or in any other country.

G protein-coupled receptors are prevalent
throughout the human body, comprising approximately 60% of
30 known cellular receptor types, and mediate signal
transduction across the cell membrane for a very wide
range of endogenous ligands. They participate in a
diverse array of physiological and pathophysiological
processes, including, but not limited to those associated
35 with cardiovascular, central and peripheral nervous
system, reproductive, metabolic, digestive, immunological,
inflammatory, and growth disorders, as well as other cell-

regulatory and proliferative disorders. Agents which selectively modulate functions of G protein-coupled receptors have important therapeutic applications. These receptors are becoming increasingly recognised as
5 important drug targets, due to their crucial roles in signal transduction (G protein-coupled Receptors, IBC Biomedical Library Series, 1996).

One of the most intensively studied G protein-coupled receptors is the receptor for C5a. C5a is one of
10 the most potent chemotactic agents known, and recruits neutrophils and macrophages to sites of injury, alters their morphology; induces degranulation; increases calcium mobilisation, vascular permeability (oedema) and neutrophil adhesiveness; contracts smooth muscle;
15 stimulates release of inflammatory mediators, including histamine, TNF- α , IL-1, IL-6, IL-8, prostaglandins, and leukotrienes, and of lysosomal enzymes; promotes formation of oxygen radicals; and enhances antibody production (Gerard and Gerard, 1994).

Agents which limit the pro-inflammatory actions of C5a have potential for inhibiting chronic inflammation, and its accompanying pain and tissue damage. For these reasons, molecules which prevent C5a from binding to its receptors are useful for treating chronic inflammatory
20 disorders driven by complement activation. Such compounds also provide valuable new insights into the mechanisms of complement-mediated immunity.
25

In our previous applications No. PCT/AU98/00490 and Australian provisional No. PR8334, the entire
30 disclosures of which are incorporated herein by this reference, we described the three-dimensional structure of some analogues of the C-terminus of human C5a, and used this information to design novel compounds which bind to the human C5a receptor (C5aR), behaving as either agonists
35 or antagonists of C5a. It had previously been thought that a putative antagonist might require both a C-terminal arginine and a C-terminal carboxylate for receptor binding

and antagonist activity (Kontetis et al, 1994). In PCT/AU98/00490 we showed that in fact a terminal carboxylate group is not generally required either for high affinity binding to C5aR or for antagonist activity.

5 Instead we found that a hitherto unrecognised structural feature, a turn conformation, was the key recognition feature for high affinity binding to the human C5a receptor on neutrophils. As described in our Australian provisional application No. PR8334, filed on 17th October 10 2001, we used these findings to design constrained structural templates which enable hydrophobic groups to be assembled into a hydrophobic array for interaction with a C5a receptor. We have subsequently found that a preferred compound of this class is able to inhibit cardiac 15 fibrosis, and this is described in our Australian provisional application No. PS1606, filed on 8th April 2002. The entire disclosures of these specifications are incorporated herein by this reference.

Asthma is a potentially life-threatening 20 condition characterized by paroxysmal attacks of bronchospasm, which cause wheezing, tightness in the chest, and difficulty in breathing. Asthmatic attacks may be provoked by exposure to a variety of stimuli, such as allergens, infection, exercise, changes in ambient 25 temperature or humidity, cigarette smoke, stress or emotional upset. Although attacks can occur at any stage of life, asthma usually has its onset during childhood; it may be associated with other hypersensitivity conditions such as eczema or hay fever. Asthma affects up to 16 30 million Americans, including approximately 10-12% of children under age 18, and its incidence is increasing. It is more common in individuals under the age of 40, and is the leading cause of absence from school and admission to hospital among children. People who have a family history 35 of asthma or who have allergies have an increased risk of developing the disease.

Treatment is predominantly with bronchodilators

such as β_2 -adrenergic agonists, administered orally or as inhaled aerosols, and corticosteroids are used in severe cases. Metered-dose inhalers, dry powder inhalers and nebulizers are widely used for inhalation therapy.

5 However, despite intensive research the available therapies are unable to control symptoms in all patients. Asthma is associated with very significant morbidity, is responsible for a high proportion of hospital emergency room admissions, and causes a number of deaths each year.
10 In addition to this, the economic cost of asthma is very high.

Eczema or eczematous dermatitis is an inflammatory dermatitis associated with itching and vesicle formation, followed by weeping and crusting of the
15 lesions. In the more chronic forms there may be lichenification and/or thickening, excoriation, or pigmentation changes; these can be disfiguring. The allergic or atopic form of eczema can be caused by a variety of allergens, which may be inhaled, contact or
20 food allergens; it can be very difficult to identify the allergen responsible. Atopic eczema is common in childhood. Topical steroids are the primary first-line treatment. Hand dermatitis is a form of atopic dermatitis, which affects an estimated 1.9 million people
25 in the United States. Chronic hand dermatitis will repeatedly relapse or flare. It can involve 25-90% of the hands and affect one or both surfaces. Treatment is usually with topical corticosteroids, but this may have limited success, and is associated with side effects such
30 as skin atrophy. Targretin, a vitamin A analogue, has recently been approved in the United States for treatment of hand dermatitis.

Dermatitis is also a major veterinary problem, and is common in companion animals such as horses, cats
35 and dogs. For example, approximately 5% to 10% of the current U.S. dog population, or four to seven million dogs, is affected. Due to the chronic nature of allergic

dermatitis in affected dogs, the estimated number of treatments per year is in excess of 8 million. Current treatment primarily uses medicated shampoos and conditioners with severe cases requiring treatment with glucocorticoids.

A variety of agents, including immunoglobulin E (IgE)-targeting monoclonal antibodies, tumour necrosis factor antagonists, immunosuppressive agent such as tacrolimus and pimecrolimus, and phototherapies, are in various stages of clinical trial for the treatment of hypersensitivity conditions.

However, there is a great need in the art for effective, non-toxic agents for the treatment of asthma and other hypersensitivity conditions, which do not require administration by injection, and which can be produced at reasonable cost. To our knowledge none of these approved or experimental agents, and in particular no small molecule agent, targets the C5a receptor.

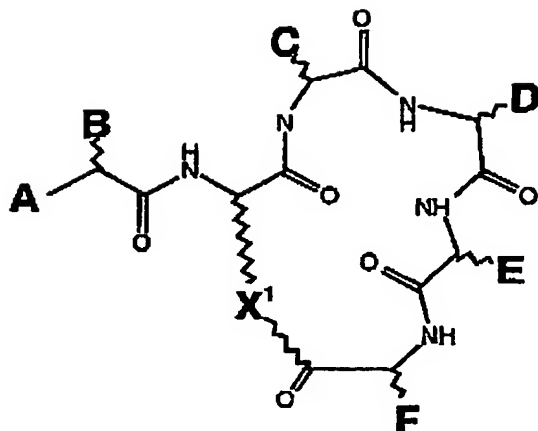
We now show for the first time that a specific inhibitor of the C5a receptor is able to ameliorate asthma in a mammal. This is the first reported case of an inhibitor of the complement system being used to modulate pathology in asthma.

SUMMARY OF THE INVENTION

According to a first aspect, the invention provides a method of treatment of a hypersensitivity conditions, comprising the step of administering an effective amount of an inhibitor of a G protein-coupled receptor to a subject in need of such treatment.

Preferably the inhibitor is a compound which

- (a) is an antagonist of a G protein-coupled receptor,
- (b) has substantially no agonist activity, and
- (c) is a cyclic peptide or peptidomimetic compound of formula I



where A is H, alkyl, aryl, NH₂, NH-alkyl,
5 N(alkyl)₂, NH-aryl, NH-acyl, NH-benzoyl, NHSO₃, NHSO₂-
alkyl, NHSO₂-aryl, OH, O-alkyl, or O-aryl;

B is an alkyl, aryl, phenyl, benzyl, naphthyl or
indole group, or the side chain of a D- or L-amino acid
such as L-phenylalanine or L-phenylglycine, but is not the
10 side chain of glycine, D-phenylalanine, L-
homophenylalanine, L-tryptophan, L-homotryptophan, L-
tyrosine, or L-homotyrosine;

C is a small substituent, such as the side chain
of a D-, L- or homo-amino acid such as glycine, alanine,
15 leucine, valine, proline, hydroxyproline, or thioproline,
but is preferably not a bulky substituent such as
isoleucine, phenylalanine, or cyclohexylalanine;

D is the side chain of a neutral D-amino acid
such as D-Leucine, D-homoleucine, D-cyclohexylalanine, D-
20 homocyclohexylalanine, D-valine, D-norleucine, D-homo-
norleucine, D-phenylalanine, D-tetrahydroisoquinoline, D-
glutamine, D-glutamate, or D-tyrosine, but is preferably
not a small substituent such as the side chain of glycine
or D-alanine, a bulky planar side chain such as D-
25 tryptophan, or a bulky charged side chain such as D-
arginine or D-Lysine;

E is a bulky substituent, such as the side chain
of an amino acid selected from the group consisting of L-

phenylalanine, L-tryptophan and L-homotryptophan, or is L-1-naphthyl or L-3-benzothienyl alanine, but is not the side chain of D-tryptophan, L-N-methyltryptophan, L-homophenylalanine, L-2-naphthyl L-
5 tetrahydroisoquinoline, L-cyclohexylalanine, D-leucine, L-fluorenylalanine, or L-histidine;

F is the side chain of L-arginine, L-homoarginine, L-citrulline, or L-canavanine, or a bioisostere thereof, ie. a side chain in which the
10 terminal guanidine or urea group is retained, but the carbon backbone is replaced by a group which has different structure but is such that the side chain as a whole reacts with the target protein in the same way as the parent group; and

15 X is $-(CH_2)_nNH-$ or $(CH_2)_nS-$, where n is an integer of from 1 to 4, preferably 2 or 3; $-(CH_2)_2O-$; $-(CH_2)_3O-$; $-(CH_2)_3-$; $-(CH_2)_4-$; $-CH_2COCHRNH-$; or $-CH_2-CHCOCHRNH-$, where R is the side chain of any common or uncommon amino acid.

20 In C, both the cis and trans forms of hydroxyproline and thioproline may be used.

Preferably A is an acetamide group, an aminomethyl group, or a substituted or unsubstituted sulphonamide group.

25 Preferably where A is a substituted sulphonamide, the substituent is an alkyl chain of 1 to 6, preferably 1 to 4 carbon atoms, or a phenyl or toluyyl group.

In a particularly preferred embodiment, the compound has antagonist activity against C5aR, and has no
30 C5a agonist activity.

The compound is preferably an antagonist of C5a receptors on human and mammalian cells including, but not limited to, human polymorphonuclear leukocytes and human macrophages. The compound preferably binds potently and
35 selectively to C5a receptors, and more preferably has potent antagonist activity at sub-micromolar concentrations. Even more preferably the compound has a

receptor affinity $IC_{50} < 25 \mu M$, and an antagonist potency $IC_{50} < 1 \mu M$.

Most preferably the compound is selected from the group consisting of compounds 1 to 6, 10 to 15, 17, 19, 20, 22, 25, 26, 28, 30, 31, 33 to 37, 39 to 45, 47 to 50, 52 to 58 and 60 to 70 described in provisional application No. PR8334. In a particularly preferred embodiment, the compound is PMX53 (compound 1), compound 33, compound 60 or compound 45 described therein.

The hypersensitivity condition may be any state in which complement-mediated tissue damage results from the immune reaction of a sensitised or immunized individual to a subsequent exposure to antigen. These include but are not limited to Type II immediate hypersensitivity (cytotoxic) and Type III (complex-mediated) immediate hypersensitivity, and Arthus-type reactions such as serum sickness, glomerulonephritis, hypereosinophilia syndrome, and farmer's lung. In one preferred embodiment the hypersensitivity condition is asthma or eczema.

The inhibitor may be used in conjunction with one or more other agents for the treatment of hypersensitivity conditions. For asthma, these include but are not limited to bronchodilators such as β_2 -adrenergic agonists, including but not limited to albuterol (Ventolin), antihistamines such as chlorpheniramine, diphenhydramine, or mepyramine, mast cell stabilisers like nedocromil, leukotriene blockers like zileutin, montelukast and zafirlukast, muscarinic antagonists like ipratropium and corticosteroids such as prednisolone, budesonide and fluticasone and synthetic steroids or analogues thereof. For eczema these include but are not limited to topical corticosteroids and synthetic steroids or analogues thereof, and Vitamin A analogues such as bexarotene ("Targretin"). The inhibitor may alternatively or additionally be used in conjunction with antiinflammatories, such as eicosapentanoic acid

derivatives and omega 3 oils.

The compounds of the invention may be formulated for oral, parenteral, inhalational, intranasal, topical or or transdermal use, but oral formulations are preferred.

5 It is expected that most if not all compounds of the invention will be stable in the presence of metabolic enzymes, such as those of the gut, blood, lung or intracellular enzymes. Such stability can readily be tested by routine methods known to those skilled in the
10 art.

Suitable formulations for administration by any desired route may be prepared by standard methods, for example by reference to well-known textbooks such as Remington: The Science and Practice of Pharmacy, Vol. II,
15 2000 (20th edition), A.R. Gennaro (ed), Williams & Wilkins, Pennsylvania.

While the invention is not in any way restricted to the treatment of any particular animal or species, it is particularly contemplated that the method of the
20 invention will be useful in medical treatment of humans, and will also be useful in veterinary treatment, particularly of companion animals such as cats and dogs, livestock such as cattle, horses and sheep, and zoo animals, including non-human primates, large bovids,
25 felids, ungulates and canids.

The compound may be administered at any suitable dose and by any suitable route. Oral, transdermal or intranasal administration is preferred, because of the greater convenience and acceptability of these routes.
30 The effective dose will depend on the nature of the condition to be treated, and the age, weight, and underlying state of health of the individual treatment. This will be at the discretion of the attending physician or veterinarian. Suitable dosage levels may readily be
35 determined by trial and error experimentation, using methods which are well known in the art.

BRIEF DESCRIPTION OF THE FIGURES

Figure 1 shows the inhibition of the vascular leakage associated with a dermal Arthus reaction by intravenous (A), oral (B) and topical (C) AcF-[OPdChaWR], and appropriate controls (D).

Figure 2 shows the inhibition of the rise in circulating TNF α associated with a dermal Arthus reaction by intravenous (A), oral (B) and topical (C) AcF-[OPdChaWR], and appropriate topical controls (D).

Figure 3 shows the reduction of the pathology index associated with a dermal Arthus reaction by intravenous, oral and topical AcF-[OPdChaWR].

DETAILED DESCRIPTION OF THE INVENTION

For the purposes of this specification it will be clearly understood that the word "comprising" means "including but not limited to", and that the word "comprises" has a corresponding meaning.

As used herein, the singular forms "a", "an", and "the" include plural reference unless the context clearly dictates otherwise. Thus, for example, a reference to "an enzyme" includes a plurality of such enzymes, and a reference to "an amino acid" is a reference to one or more amino acids. Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. Although any materials and methods similar or equivalent to those described herein can be used to practice or test the present invention, the preferred materials and methods are now described.

Abbreviations used herein are as follows:

D-Cha	D-cyclohexylamine
LPS	lipopolysaccharide
PMN	polymorphonuclear granulocyte
RMSD	root mean square deviation

rp-HPLC reverse phase-high performance liquid
chromatography
TFA trifluoroacetic acid;

5 Throughout the specification conventional single-letter and three-letter codes are used to represent amino acids.

For the purposes of this specification, the term "alkyl" is to be taken to mean a straight, branched, or
10 cyclic, substituted or unsubstituted alkyl chain of 1 to 6, preferably 1 to 4 carbons. Most preferably the alkyl group is a methyl group. The term "acyl" is to be taken to mean a substituted or unsubstituted acyl of 1 to 6, preferably 1 to 4 carbon atoms. Most preferably the acyl
15 group is acetyl. The term "aryl" is to be understood to mean a substituted or unsubstituted homocyclic or heterocyclic aryl group, in which the ring preferably has 5 or 6 members.

A "common" amino acid is a L-amino acid selected
20 from the group consisting of glycine, leucine, isoleucine, valine, alanine, phenylalanine, tyrosine, tryptophan, aspartate, asparagine, glutamate, glutamine, cysteine, methionine, arginine, lysine, proline, serine, threonine and histidine.

25 An "uncommon" amino acid includes, but is not restricted to, D-amino acids, homo-amino acids, N-alkyl amino acids, dehydroamino acids, aromatic amino acids other than phenylalanine, tyrosine and tryptophan, ortho-, meta- or para-aminobenzoic acid, ornithine, citrulline,
30 canavanine, norleucine, γ -glutamic acid, aminobutyric acid, L-fluorenylalanine, L-3-benzothienylalanine, and α,α -disubstituted amino acids.

Generally, the terms "treating", "treatment" and the like are used herein to mean affecting a subject,
35 tissue or cell to obtain a desired pharmacological and/or physiological effect. The effect may be prophylactic in terms of completely or partially preventing a disease or

sign or symptom thereof, and/or may be therapeutic in terms of a partial or complete cure of a disease.

"Treating" as used herein covers any treatment of, or prevention of disease in a vertebrate, a mammal, particularly a human, and includes: preventing the disease from occurring in a subject who may be predisposed to the disease, but has not yet been diagnosed as having it; inhibiting the disease, i.e., arresting its development; or relieving or ameliorating the effects of the disease, i.e., cause regression of the effects of the disease.

The invention includes the use of various pharmaceutical compositions useful for ameliorating disease. The pharmaceutical compositions according to one embodiment of the invention are prepared by bringing a compound of formula I, analogue, derivatives or salts thereof and one or more pharmaceutically-active agents or combinations of compound of formula I and one or more pharmaceutically-active agents into a form suitable for administration to a subject using carriers, excipients and additives or auxiliaries.

Frequently used carriers or auxiliaries include magnesium carbonate, titanium dioxide, lactose, mannitol and other sugars, talc, milk protein, gelatin, starch, vitamins, cellulose and its derivatives, animal and vegetable oils, polyethylene glycols and solvents, such as sterile water, alcohols, glycerol and polyhydric alcohols. Intravenous vehicles include fluid and nutrient replenishers. Preservatives include antimicrobial, anti-oxidants, chelating agents and inert gases. Other pharmaceutically acceptable carriers include aqueous solutions, non-toxic excipients, including salts; preservatives, buffers and the like, as described, for instance, in Remington's Pharmaceutical Sciences, 20th ed. Williams & Wilkins (2000) and The British National Formulary 43rd ed. (British Medical Association and Royal Pharmaceutical Society of Great Britain, 2002; <http://bnf.rhn.net>), the contents of which are hereby

incorporated by reference. The pH and exact concentration of the various components of the pharmaceutical composition are adjusted according to routine skills in the art. See Goodman and Gilman's The Pharmacological
5 Basis for Therapeutics (7th ed., 1985).

The pharmaceutical compositions are preferably prepared and administered in dosage units. Solid dosage units include tablets, capsules and suppositories. For treatment of a subject, depending on activity of the
10 compound, manner of administration, nature and severity of the disorder, age and body weight of the subject, different daily doses can be used. Under certain circumstances, however, higher or lower daily doses may be appropriate. The administration of the daily dose can be
15 carried out both by single administration in the form of an individual dose unit or else several smaller dose units and also by multiple administration of subdivided doses at specific intervals.

The pharmaceutical compositions according to the
20 invention may be administered locally or systemically in a therapeutically effective dose. Amounts effective for this use will, of course, depend on the severity of the disease and the weight and general state of the subject. Typically, dosages used *in vitro* may provide useful
25 guidance in the amounts useful for *in situ* administration of the pharmaceutical composition, and animal models may be used to determine effective dosages for treatment of the cytotoxic side effects. Various considerations are described, eg. in Langer, Science, 249: 1527, (1990).
30 Formulations for oral use may be in the form of hard gelatin capsules, in which the active ingredient is mixed with an inert solid diluent, for example, calcium carbonate, calcium phosphate or kaolin. They may also be in the form of soft gelatin capsules, in which the active
35 ingredient is mixed with water or an oil medium, such as peanut oil, liquid paraffin or olive oil.

Aqueous suspensions normally contain the active materials in admixture with excipients suitable for the manufacture of aqueous suspensions. Such excipients may be suspending agents such as sodium carboxymethyl
5 cellulose, methyl cellulose, hydroxypropylmethylcellulose, sodium alginate, polyvinylpyrrolidone, gum tragacanth and gum acacia; dispersing or wetting agents, which may be (a) a naturally occurring phosphatide such as lecithin; (b) a condensation product of an alkylene oxide with a fatty
10 acid, for example, polyoxyethylene stearate; (c) a condensation product of ethylene oxide with a long chain aliphatic alcohol, for example, heptadecaethylenoxycetanol; (d) a condensation product of ethylene oxide with a partial ester derived from a fatty
15 acid and hexitol such as polyoxyethylene sorbitol monooleate, or (e) a condensation product of ethylene oxide with a partial ester derived from fatty acids and hexitol anhydrides, for example polyoxyethylene sorbitan monooleate.

20 The pharmaceutical compositions may be in the form of a sterile injectable aqueous or oleaginous suspension. This suspension may be formulated according to known methods using suitable dispersing or wetting agents and suspending agents such as those mentioned
25 above. The sterile injectable preparation may also a sterile injectable solution or suspension in a non-toxic parenterally-acceptable diluent or solvent, for example, as a solution in 1,3-butanediol. Among the acceptable vehicles and solvents which may be employed are water,
30 Ringer's solution, and isotonic sodium chloride solution. In addition, sterile, fixed oils are conventionally employed as a solvent or suspending medium. For this purpose, any bland fixed oil may be employed, including synthetic mono-or diglycerides. In addition, fatty acids
35 such as oleic acid may be used in the preparation of injectables.

Compounds of formula I may also be administered in the form of liposome delivery systems, such as small unilamellar vesicles, large unilamellar vesicles, and multilamellar vesicles. Liposomes can be formed from a variety of phospholipids, such as cholesterol, stearylamine, or phosphatidylcholines.

Dosage levels of the compound of formula I of the present invention will usually be of the order of about 0.5mg to about 20mg per kilogram body weight, with a preferred dosage range between about 0.5mg to about 10mg per kilogram body weight per day (from about 0.5g to about 3g per patient per day). The amount of active ingredient which may be combined with the carrier materials to produce a single dosage will vary, depending upon the host to be treated and the particular mode of administration. For example, a formulation intended for oral administration to humans may contain about 5mg to 1g of an active compound with an appropriate and convenient amount of carrier material, which may vary from about 5 to 95 percent of the total composition. Dosage unit forms will generally contain between from about 5mg to 500mg of active ingredient.

It will be understood, however, that the specific dose level for any particular patient will depend upon a variety of factors including the activity of the specific compound employed, the age, body weight, general health, sex, diet, time of administration, route of administration, rate of excretion, drug combination and the severity of the particular disease undergoing therapy.

In addition, some of the compounds of the invention may form solvates with water or common organic solvents. Such solvates are encompassed within the scope of the invention.

The compounds of the invention may additionally be combined with other therapeutic compounds to provide an operative combination. It is intended to include any chemically compatible combination of pharmaceutically-

active agents, as long as the combination does not eliminate the activity of the compound of formula I of this invention.

In evaluation of the compounds of the invention, conventional measures of efficacy may be used. For example, for asthma commonly-used primary efficacy endpoints include lung function tests such as spirometry or measurement of vital capacity, or self-monitoring using a peak flow meter. For eczema, evaluation of efficacy may be based on:

- (a) Physician's Static Assessment (PSA), a primary endpoint required by the United States Food and Drug Administration, which calls for 90% or greater improvement in signs and symptoms, is equivalent to a clear or almost clear condition on at least two observations 21 days apart, or
- (b) Physician's Global Assessment (PGA), which calls for 50% or greater improvement.

The invention will now be described by way of reference only to the following general methods and experimental examples.

General Methods

Cyclic peptide compounds of formula I are prepared according to methods described in detail in our earlier applications No. PCT/AU98/00490 and PR8334, the entire disclosures of which are incorporated herein by this reference. While the invention is specifically illustrated with reference to the compound AcF-[OPdChaWR] (PMX53), whose corresponding linear peptide is Ac-Phe-Orn-Pro-dCha-Trp-Arg, it will be clearly understood that the invention is not limited to this compound.

Compounds 1-6, 17, 20, 28, 30, 31, 36 and 44 disclosed in International patent application No. PCT/AU98/00490 and compounds 10-12, 14, 15, 25, 33, 35, 40, 45, 48, 52, 58, 60, 66, and 68-70 disclosed for the first time in Australian provisional application No.

PR8334 have appreciable antagonist potency ($IC_{50} < 1 \mu M$) against the C5a receptor on human neutrophils. PMX53 (compound 17 of PCT/AU98/00490; also identified as compound 1 in PR8334) and compounds 33, 45 and 60 of PR8334 are most preferred.

We have found that all of the compounds of formula I which have so far been tested have broadly similar pharmacological activities, although the physicochemical properties, potency, and bioavailability of the individual compounds vary somewhat, depending on the specific substituents.

The following general tests may be used for initial screening of candidate inhibitor of G protein-coupled receptors, and especially of C5a receptors.

Receptor-Binding Assay

Assays were performed with fresh human PMNs, isolated as previously described (Sanderson et al, 1995), using a buffer of 50 mM HEPES, 1 mM $CaCl_2$, 5 mM $MgCl_2$, 0.5% bovine serum albumin, 0.1% bacitracin and 100 μM phenylmethylsulfonyl fluoride (PMSF). In assays performed at 4°C, buffer, unlabelled human recombinant C5a (Sigma) or peptide, Hunter/Bolton labelled ^{125}I -C5a (~ 20 pM) (New England Nuclear, MA) and PMNs (0.2×10^6) were added sequentially to a Millipore Multiscreen assay plate (HV 0.45) having a final volume of 200 μL /well. After incubation for 60 min at 4°C, the samples were filtered and the plate washed once with buffer. Filters were dried, punched and counted in an LKB gamma counter. Non-specific binding was assessed by the inclusion of 1mM peptide or 100 nM C5a, which typically resulted in 10-15% total binding.

Data was analysed using non-linear regression and statistics with Dunnett post-test.

Myeloperoxidase Release Assay for Antagonist Activity

Cells were isolated as previously described

(Sanderson et al, 1995) and incubated with cytochalasin B (5µg/mL, 15 min, 37°C). Hank's Balanced Salt solution containing 0.15% gelatin and peptide was added on to a 96 well plate (total volume 100 µL/well), followed by 5 25 µL cells (4×10^6 /mL). To assess the capacity of each peptide to antagonise C5a, cells were incubated for 5 min at 37°C with each peptide, followed by addition of C5a (100 nM) and further incubation for 5 min. Then 50 µL of sodium phosphate (0.1M, pH 6.8) was added to each well, 10 the plate was cooled to room temperature, and 25 µL of a fresh mixture of equal volumes of dimethoxybenzidine (5.7 mg/mL) and H₂O₂ (0.51%) was added to each well. The reaction was stopped at 10 min by addition of 2% sodium azide. Absorbances were measured at 450 nm in a Bioscan 15 450 plate reader, corrected for control values (no peptide), and analysed by non-linear regression.

In Vivo Assays of Anti-Inflammatory Activity

The following well-known *in vivo* assay systems 20 are used to assess the anti-inflammatory activity of compounds of the invention. All assay data are analysed using non-linear regression analysis and Student's t-test, analysis of variance, with $p < 0.05$ as the threshold level of significance.

25

(a) Carrageenan Paw Oedema

Anaesthetised (i.p. ketamine & xylazine) Wistar rats (150-200g) or mice are injected with sterilised air (20ml day 1, 10ml day 4) into the subcutaneous tissue of 30 the back. The cavity can be used after 6 days, whereupon carrageenan (2ml, 1% w/w in 0.9% saline) is injected into the air pouch, and exudate is collected after 10 hr. Test compounds are administered daily after Day 6, and their anti-inflammatory effects assayed by differential counting 35 of cells in the air-pouch exudate. Animals are killed at appropriate times after injection, and 2ml 0.9% saline is used to lavage the cavity; lavage fluids are transferred

to heparinised tube and cells are counted with a haemocytometer and Diff-Quik stained cytocentrifuged preparation.

Alternatively, a routine carrageenan paw oedema developed in Wistar rats by administering a pedal injection of carrageenan may be used to elicit oedema which is visible in 2h and maximised in 4h. Test compounds are given 40 min before inflammatory stimulus and evaluated by microcaliper measurements of paws after 2 & 4 hr. See Fairlie, D.P. et al (1987). Also see Walker and Whitehouse (1978).

(b) Adjuvant Arthritis.

Adjuvant arthritis is induced in rats (3 strains) either microbially (injection of heat-killed *Mycobacterium tuberculosis*) or chemically (with avridine) by inoculation with the arthritogenic adjuvant co-administered with oily vehicles (Freund's adjuvants) in the tail base. (See Whitehouse, M. W., Handbook of Animal Models for the Rheumatic Diseases, Eds. Greenwald, R. A.; Diamond, H. S.; Vol. 1, pp. 3-16, CRC Press)

Within 13 days the adjuvant arthritis is manifested by local inflammation and ulceration in the tail, gross swelling of all four paws, inflammatory lesions in paws and ears, weight loss and fever. These symptoms, which are similar to those of inflammatory disease in humans (Winter and Nuss, 1966), can be alleviated by agents such as indomethacin or cyclosporin, which also show beneficial effects in man (eg. Ward and Cloud, 1966). Without drug treatment at Day 14, arthritic rats had hypertrophy of the paws, reduced albumin but raised acute phase reaction proteins in serum, and depressed hepatic metabolism of xenobiotics as indicated by prolonged barbiturate-induced sleeping times.

To assess activity, compounds are administered for 4 days orally ($\leq 10\text{mg/kg/day}$) or (intraperitoneally (i.p.) from Days 10-13 following inoculation with

arthritogen (Day 0). If the compound is active, the inflammation is either not visible, or is very significantly reduced in rear or front paws, as assessed by microcaliper measurements of paw thickness and tail volume, as well as by gross inspection of inflammatory lesions. Animals are sacrificed by cervical dislocation on Day 18 unless arthritis signs are absent, whereupon duration of observations is continued with special permission from the Ethics Committees. Experiments are staggered to maximise throughput and allow early comparisons between compounds. This routine assay is well-accepted as identifying anti-inflammatory agents for use in humans.

15 Example 1 Reverse Passive Arthus Reaction in the Rat

A reverse passive peritoneal Arthus reaction was induced as previously described (Strachan et al., 2000), and a group of rats were pretreated prior to peritoneal deposition of antibody with AcF-[OPdChawR] (1) by oral gavage (10mg kg⁻¹ dissolved in 10% ethanol/90% saline solution to a final volume of 200µl) or an appropriate oral vehicle control 30 min prior to deposition of antibody. Female Wistar rats (150-250g) were anaesthetised with ketamine (80mg kg⁻¹ i.p.) and xylazine (12mg kg⁻¹ i.p.).

25 The lateral surfaces of the rat were carefully shaved and 5 distinct sites on each lateral surface clearly delineated. A reverse passive Arthus reaction was induced in each dermal site by injecting Evans blue (15 mg kg⁻¹ i.v.), chicken ovalbumin (20mg kg⁻¹ i.v.) into the femoral vein 10min prior to the injection of antibody. Rabbit anti-chicken ovalbumin (saline only, 100, 200, 300 or 400µg antibody in a final injection volume of 30µL) was injected in duplicate at two separate dermal sites on each lateral surface of the rat, giving a total of 10 injection sites per rat. Rats were placed on a heating pad, and anaesthetic was maintained over a 4h-treatment period with periodic collection of blood samples. Blood was allowed

to spontaneously clot on ice, and serum samples were collected and stored at -20°C . Four hours after induction of the dermal Arthus reaction, the anaesthetised rat was euthanased and a 10mm^2 area of skin was collected from the site of each Arthus reaction. Skin samples were stored in 10% buffered formalin for at least 10 days before histological analysis using haematoxylin and eosin stain. Additionally, a second set of skin samples were placed in 1mL of formamide overnight, and the absorbance of Evans blue extraction measured at 650nm, as an indicator of serum leakage into the dermis. Figure 1 shows the optical density of dermal punch extracts following intradermal injection of rabbit anti-chicken ovalbumin at $0-400\text{ }\mu\text{g site}^{-1}$ following pretreatment with AcF-[OPdChaWR] intravenously, orally or topically. Data are shown as absorbance at 650nm as a percentage of the plasma absorbance, as mean values \pm SEM ($n=3-6$). *indicates a P value ≤ 0.05 when compared to Arthus control values.

Rats were pretreated with the C5aR antagonist, AcF-[OPdChaWR] (1) as the TFA salt, either intravenously ($0.3-1\text{mg kg}^{-1}$ in $200\mu\text{L}$ saline containing 10% ethanol, 10min prior to initiation of dermal Arthus), orally ($0.3-10\text{mg kg}^{-1}$ in $200\text{ }\mu\text{L}$ saline containing 10% ethanol by oral gavage, 30min prior to initiation of dermal Arthus in rats denied food access for the preceding 18hours) or topically ($200-400\mu\text{g site}^{-1}$ 10min prior to initiation of dermal Arthus reaction), or with the appropriate vehicle control. Topical application of the antagonist involved application of $20\mu\text{L}$ of a $10-20\text{mg mL}^{-1}$ solution in 10% dimethyl sulphoxide (DMSO), which was then smeared directly onto the skin at each site, 10min prior to induction of the Arthus reaction.

The saline-only injection site from rats treated with Evans blue only served as antigen controls, the saline-only injection site from rats treated with Evans blue plus topical DMSO only served as a vehicle control, the saline-only injection site from rats treated with

Evans blue plus either intravenous, oral or topical antagonist only served as antagonist controls, and Evans blue plus dermal rabbit anti-chicken ovalbumin served as antibody controls. Topical application of the peptide
5 AcF-[OPGWR] which has similar chemical composition and solubility as AcF-[OPdChawR] (1), but with an IC_{50} binding affinity of $>1mM$ in isolated human PMNs, served as an inactive peptide control. AcF-[OPGWR] was also dissolved in 10% DMSO and applied topically at $400\mu g$ site⁻¹ 10 min
10 prior to initiation of the Arthus reaction.

TNF α Measurement

Serum TNF α concentrations were measured using an enzyme-linked immunosorbent assay (ELISA) kit (Strachan et
15 al., 2000). Antibody pairs used were a rabbit anti-rat TNF α antibody coupled with a biotinylated murine anti-rat TNF α antibody. Figure 2 shows the serum TNF α concentrations at regular intervals after initiation of a dermal Arthus reaction, with group of rats pretreated with
20 AcF-[OPdChawR] intravenously, orally or topically. Data are shown as mean values \pm SEM ($n=3-6$). *indicates a P value of ≤ 0.05 when compared to Arthus control values.

Interleukin-6 Measurement

25 An ELISA method as described previously was used to measure serum and peritoneal lavage fluid interleukin-6 (IL-6) concentrations (Strachan et al., 2000).

Pathology Assessment

30 Rat skin samples were fixed in 10% buffered formalin for at least 10 days, and stained with haematoxylin and eosin using standard histological techniques. Dermal samples were analysed in a blind fashion for evidence of pathology, and the degree of rat
35 PMN infiltration was scored on a scale of 0-4. Initiation of a dermal Arthus reaction resulted in an increase in interstitial neutrophils, which was quantified in the

following manner. Sections were given a score of 0 if no abnormalities were detected. A score of 1 indicated the appearance of increased PMNs in blood vessels, but no migration of inflammatory cells out of the lumen. A score of 2 and 3 indicated the appearance of increasing numbers of PMNs in the interstitial tissue and more prominent accumulations of inflammatory cells around blood vessels. A maximal score of 4 indicated severe pathological abnormalities were present in dermal sections, with excessive infiltration of PMNs into the tissues and migration of these cells away from blood vessels. Figure 3 shows the intradermal injection of increasing amounts of antibody leads to a dose-responsive increase in the pathology index scored by dermal samples (A). Data are shown for dermal samples intradermally injected with saline or 400 $\mu\text{g site}^{-1}$ antibody ($n=5$) in rats pretreated with AcF-[OPdChaWR] intravenously (B) ($n=3$), orally (C) ($n=3$) and topically (D) ($n=3$). Data are shown as mean values \pm SEM. * $P \leq 0.05$ when compared to Arthus values using a non-parametric t-test.

Example 3 Treatment of Asthma in a Tiger Cub

Respiratory problems were first noticed in Kaasha, a female Bengal Tiger cub at the Dreamworld park, Australia, at the age of 10 weeks and around 10kg body weight. The initial clinical signs observed were a mild to moderate increase in respiratory effort after feeding, followed within a few days by a continuous increase in respiratory effort. At no time prior to the initial clinical signs had Kaasha's keepers noticed any change in demeanour, appetite, activity level, or other parameter that might indicate illness. The rectal temperature was normal at the time of initial veterinary examination, and remained normal when measured over the following weeks. The principal clinical signs included increased respiratory effort characterised by a prolonged two-phase forced

expiration, fine pulmonary crackles particularly in dorsal lobes, and a bronchointerstitial pattern and air entrapment on radiographs.

Initial treatment, pending definitive diagnosis, included antibiotics, Clavulox (7 mg/kg administered subcutaneously), doxycycline (5mg/kg administered orally), and enrofloxacin (5mg/kg administered subcutaneously), and terbutaline (0.3 mg/kg administered orally).

Two weeks after initial detection of clinical signs, Kaasha experienced an episode of severe dyspnoea with open-mouthed breathing. The episode lasted 20 to 30 seconds and resolved spontaneously.

Following repeated clinical examinations, radiography, clinical pathology (blood: CBC, MBA) (Blood biochemistry and haematology) and bronchoalveolar lavage cytology, a diagnosis of lower airway inflammation, of unknown aetiology, was made. Culture of bronchoalveolar lavage samples for microorganisms did not indicate the presence of any bacterial infection.

The anatomical diagnosis of neutrophil-dominated lower airway inflammation was confirmed histologically by a thoracoscopically-guided lung biopsy, although the pathologist also noted mixed inflammation of the interstitium. Further culture of the biopsy and polymerase chain reaction (PCR) for Feline viral rhinotracheitis virus and Feline Calicivirus and *Chlamydia* failed to provide convincing information regarding the aetiology of the condition.

Approximately three weeks after initial clinical signs, the treatment included Clavulox (7 mg/kg administered subcutaneously), doxycycline (5mg/kg administered orally), and enrofloxacin (5mg/kg administered subcutaneously) and terbutaline (0.3 mg/kg administered orally), nebulisation (saline) and percussion, use of Ventolin 100µg by puffer inhalation and terbutaline (0.3 mg/kg administered orally) as necessary to control dyspnoea. Prednisolone was given as a single daily dose of

2mg/kg, and Seretide (Salmeterol 50µg plus fluticasone 250 µg) was given using a mask and spacer. Response to treatment was marked, with improvement in respiratory effort and reduction in crackles audible on auscultation.

5 This combination of treatment was maintained over the next month.

During this month of therapy the following observations were made:

- 10 1. There was a radiographic improvement characterised by reduction in the prominence of the bronchointerstitial pattern and reduction in air entrapment.
2. Throughout each day, and from day to day, there was a marked variation in severity of respiratory clinical signs, although these were not as severe as those
- 15 observed in the initial stages of the disease. Increase in respiratory effort was often observed when the cub was taken into the airconditioned nursery, during exercise or stress, and spontaneously, presumably in response to respiratory irritants in the environment.
- 20 3. Respiratory clinical signs responded rapidly to bronchodilators, given either orally terbutaline (0.3 mg/kg orally) or by inhalation (Ventolin puffer 100µg).
4. Although response to therapy was marked, there were never times when the breathing pattern was normal.
- 25 5. The cub developed a poor "staring" coat, poor muscling, retarded growth rate, reduced activity level and playfulness, and relatively poor appetite when compared with her littermate.

The clinical findings and response to various

30 therapies were reviewed, and the diagnosis of feline asthma was made. This decision was based upon the marked reactivity of the airways, marked and rapid response to bronchodilators, and general improvement of respiratory clinical signs with corticosteroid therapy.

35 Three months after onset of clinical signs, despite relatively aggressive oral corticosteroid therapy Prednisolone 2mg/kg once daily, the level of control of

the disease was stable but not yet satisfactory in terms of long-term health management. Consequently an experimental technique involving direct injection of Prednisolone sodium succinate, 1200 mg in a total volume of 30 ml (40mg/ml aqueous solution), was instilled into the trachea under general anaesthesia. Recovery from the anaesthesia was uneventful, and within 24 hours there was a rapid and marked improvement in respiratory effort.

Kaasha's keepers unanimously reported that the breathing pattern improved to virtually normal levels for one week, with no episodes of dyspnoea during that period. However, the clinical pattern of laboured breathing returned to pre-treatment levels at seven to nine days post treatment. At this time therapy included oral corticosteroid at 2mg/kg, use of inhaled Flixotide (Fluticasone 250µg/dose) with and without Seretide (Salmeterol 50µg plus fluticasone 250µg / dose), use of Ventolin puffer, 100µg, as necessary to control dyspnoea, and occasional use of Pulmicort nebules (Budesonide 400µg) by nebulisation.

It is important to note that use of inhaled medications in the cub was characterised by variability in the effectiveness of the daily dose given, as a result of variation in her compliance, keeper compliance, keeper competence, and daily frequency of administration.

Three weeks after the intra-airway steroid procedure, another experimental therapy was applied. Injections of the C5a complement receptor antagonist AcF-[OpdChawR] (1) were administered at a dose rate of 0.3mg/kg as single daily subcutaneous injections for six days, followed by twice-weekly injections for 8 weeks. At this time the dose of oral corticosteroid, Prednisolone approx 1mg/kg reducing, had been reduced to 20mg per day because of concern regarding side effects of prolonged high dose use. Kaasha's keepers were asked to pay particular attention to ensuring that inhaled medications

were being applied in the most effective manner, to compensate for reduction in the oral corticosteroid dose.

There was unanimous agreement among Kaasha's keepers that there was a moderate to marked improvement in breathing and in recovery time after episodes of dyspnoea following the week of daily AcF-[OpdChaWR] (1) injections. However, it was observed that the general breathing pattern was not as good during twice weekly treatments as it had been following daily injections, although recovery time was comparable.

The reduction in oral cortisone dosage and treatment with AcF-[OPdChaWR] (1) corresponded with a marked improvement in the playfulness, general activity level, appetite and general demeanour of the cub. The addition of a further medication, Singulaire, has not been associated with a noticeable improvement in clinical signs.

As at July 2002 Kaasha was approximately 32 kg and 7 months of age, and was being maintained on the following regimen:

Macrolone (prednisolone): 20mg orally each evening;
Singulaire (montelukast sodium): 10mg orally each evening;
AcF-[OpdChaWR] (1): 3mg/10kg subcutaneously twice a week;
Seretide puffer (Salmeterol 50µg plus fluticasone 250µg / dose:) morning and night, preceded by Ventolin, 100µg, puffer;
Flixotide puffer (Fluticasone): 250µg/dose up to four times each day; and
Pulmicort nebulisation (Budesonide): 400µg once a day as time permits.

Further improvements in the therapeutic regime were tested with a view to long-term control of the asthma. Measures were taken to improve the efficiency of puffer medication delivery in an attempt to reduce or eliminate the oral corticosteroid use. At this time Kaasha showed mild dyspnoea at rest, and moderate to marked dyspnoea after exercise, exertion or stress, but

this dyspnoea was not associated with visible distress. Her behaviour, growth rate and appetite were only slightly less than, or comparable with, her those of littermate.

5 A dietary trial was attempted, with complete replacement of the current diet by a different protein source, namely either lamb/mutton or rabbit exclusively.

10 Kaasha's condition deteriorated, and in early September 2002 she died under anaesthetic while undergoing a brain scan. From the pathology reports it appears that the hyperoesinophilia was affecting other organs apart from the lung and the animal was becoming increasingly ill from intestinal and renal effects. While it is not possible to draw any causal connection, it is noted that in August 2002 treatment with PMX53 had been discontinued.
15 Post-mortem examination showed that the tiger cub had hyperoesinophilia syndrome, which contributed to the asthma-like lung condition, and also caused lesions in the kidney and intestine. This condition also occurs in humans.

20

DISCUSSION

Cyclic peptides have several important advantages over acyclic peptides as drug candidates (Fairlie et al 1995, Fairlie et al, 1998, Tyndall and Fairlie, 2001).
25 The cyclic compounds described in this specification are stable to proteolytic degradation for at least several hours at 37°C in human blood or plasma, in human or rat gastric juices, or in the presence of digestive enzymes such as pepsin, trypsin and chymotrypsin. In contrast,
30 short linear peptides composed of L-amino acids are rapidly degraded to their component amino acids within a few minutes under these conditions. A second advantage lies in the 'constrained single conformations adopted by the cyclic and non-peptidic molecules, in contrast to
35 acyclic or linear peptides, which are flexible enough to adopt multiple structures in solution other than the one required for receptor-binding. Thirdly, cyclic compounds

such as those described in this invention are usually more lipid-soluble and more pharmacologically bioavailable as drugs than acyclic peptides, which can rarely be administered orally. Fourthly, the plasma half-lives of cyclic molecules are usually longer than those of peptides.

It will be apparent to the person skilled in the art that while the invention has been described in some detail for the purposes of clarity and understanding, various modifications and alterations to the embodiments and methods described herein may be made without departing from the scope of the inventive concept disclosed in this specification.

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FIGURE 1

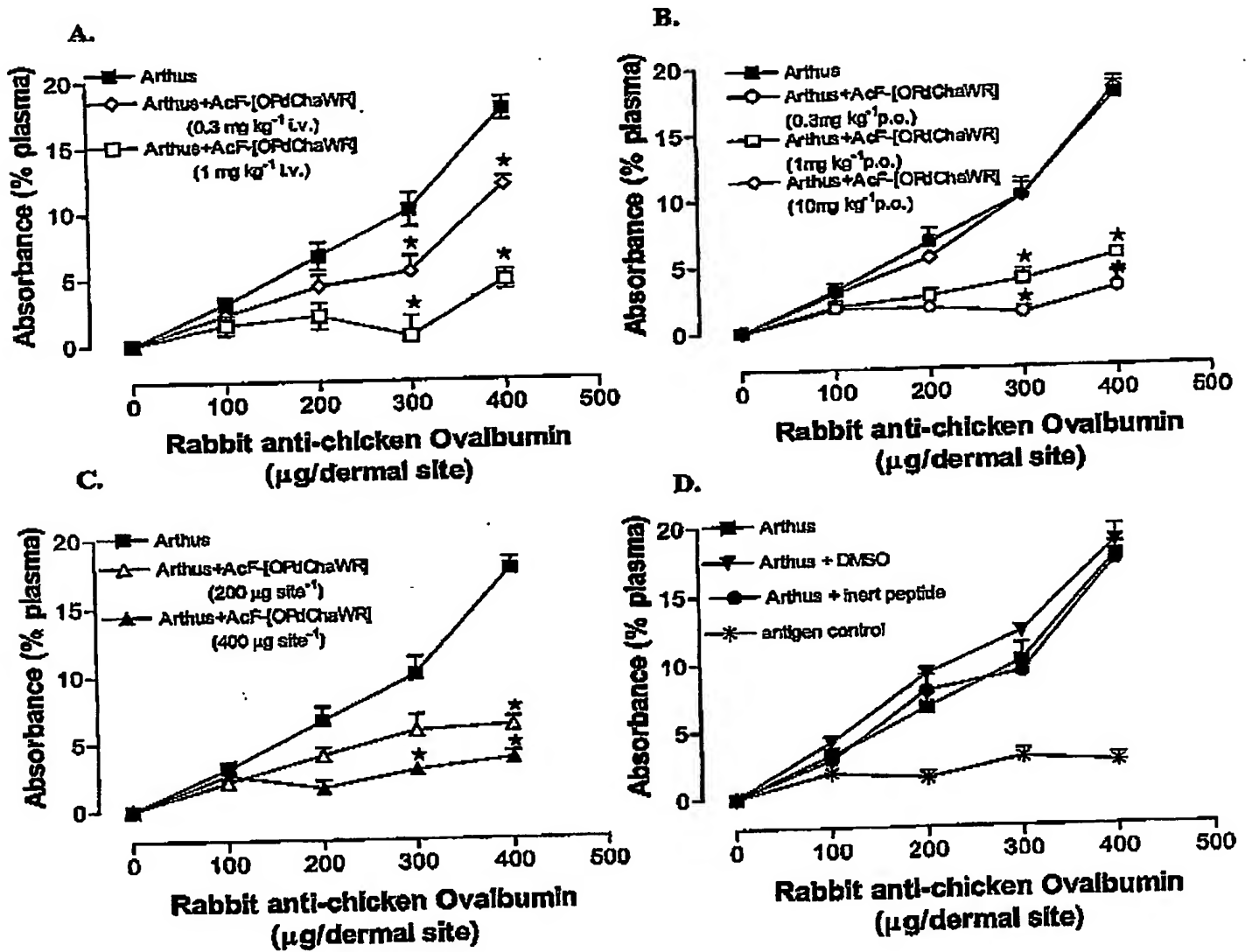


FIGURE 2

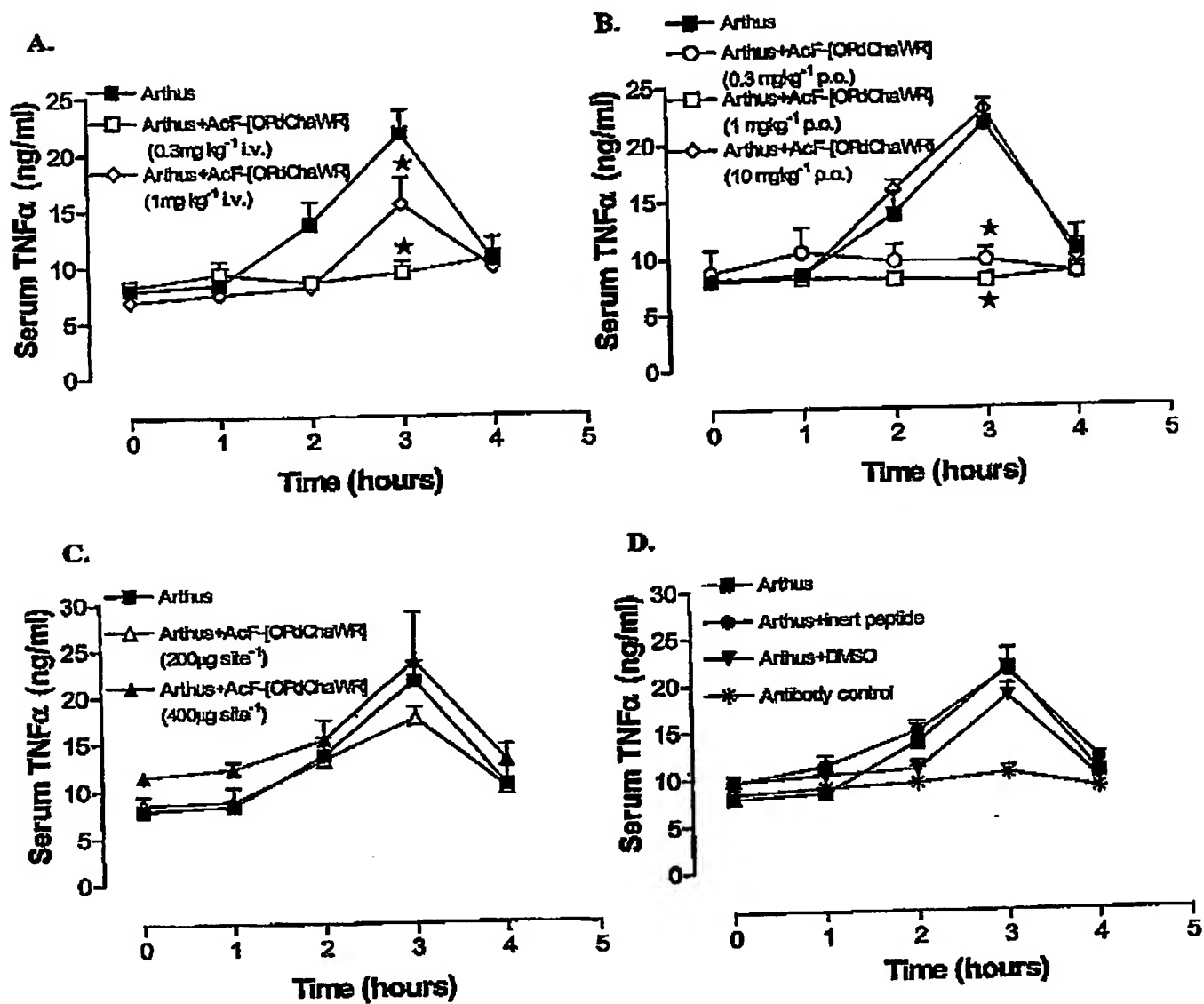
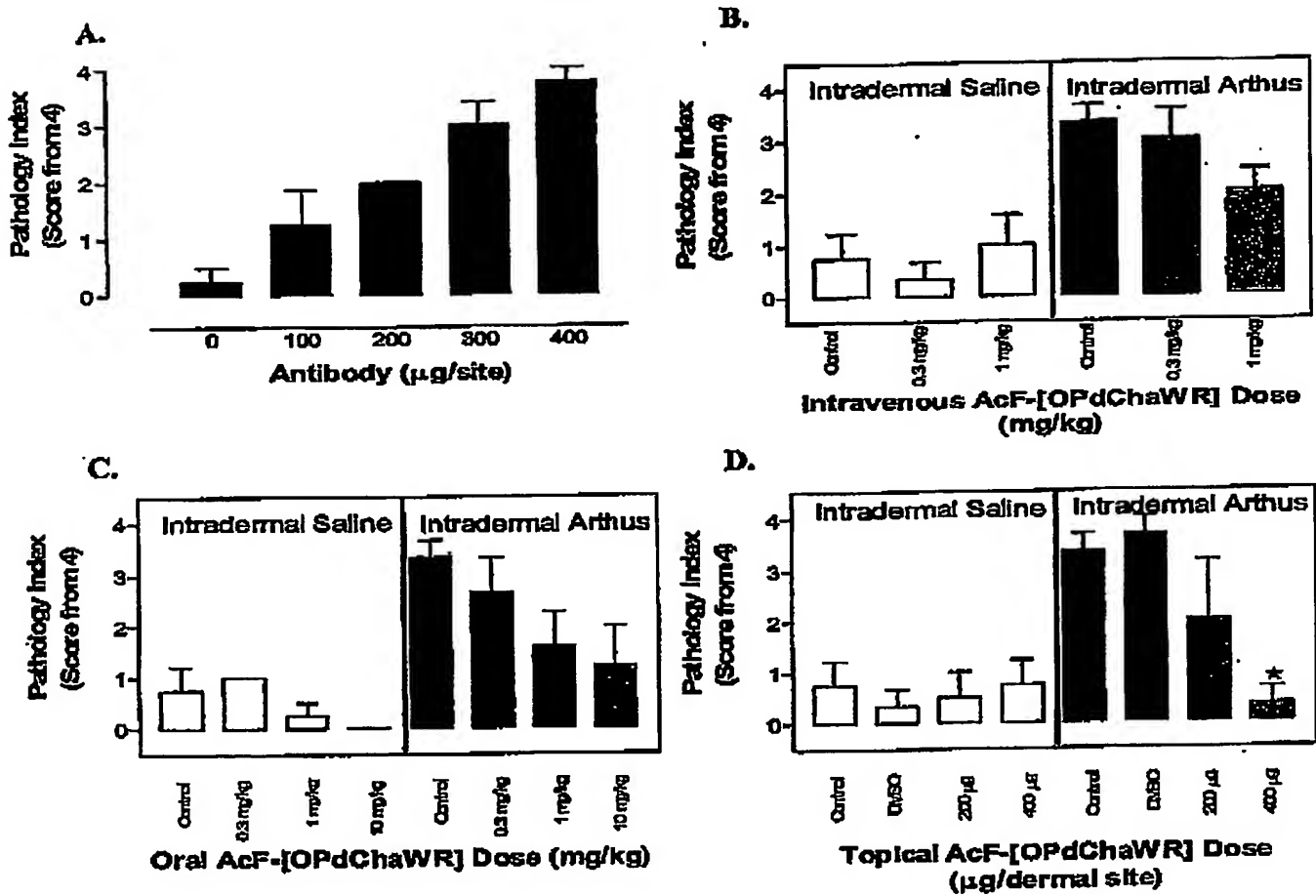


FIGURE 3



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